

Reconsideration and allowance are respectfully requested.

### AMENDMENT

Please amend the claims as follows:

a  
1. (Amended) A glycosylated or nonglycosylated protein having agonist and/or antagonist activity of the formula

$\beta^1$ -(linker<sup>1</sup>)<sub>m</sub>- $\alpha$ -(linker<sup>2</sup>)<sub>n</sub>- $\beta^2$  (1); or

$\beta^1$ -(linker<sup>1</sup>)<sub>m</sub>- $\beta^2$ -(linker<sup>2</sup>)<sub>n</sub>- $\alpha$  (2); or

$\alpha$ -(linker<sup>1</sup>)<sub>m</sub>- $\beta^1$ -(linker<sup>2</sup>)<sub>n</sub>- $\beta^2$  (3)

wherein each of  $\beta^1$  and  $\beta^2$  has the amino acid sequence of the  $\beta$  subunit of a vertebrate glycoprotein hormone which is selected from the group consisting of thyroid stimulating hormone (TSH), follicle stimulating hormone (FSH), leutinizing hormone (LH) and chorionic gonadotrophin (CG) or a variant thereof which variant binds to the receptor for said  $\beta$ -subunit;

" $\alpha$ " designates the  $\alpha$  subunit of a vertebrate glycoprotein hormone TSH, FSH, LH or CG or a variant thereof;

"linker" refers to a covalently linked moiety that spaces the  $\beta^1$  and  $\beta^2$  subunits at distances from the  $\alpha$  subunit and from each other effective to retain said activity, and

each of m and n is independently 0 or 1;

wherein said agonist and/or antagonist activity is with respect to the receptor for which at least one of said  $\beta$  subunits is a ligand.

Please cancel claim 3.

b  
2. (Amended) The protein of claim 1 wherein [the  $\alpha$  subunit or] one or more of the  $\alpha$  and  $\beta$  subunits [or both] are modified by the insertion of a CTP unit or variant thereof into a noncritical region thereof and/or wherein said linker moiety includes a CTP unit or variant thereof.

5. (Amended) The protein of claim 1 wherein said variants contain 1-5 conservative amino acid substitutions as referred to the native forms or [are truncated forms] lack 1-10 amino acids at the N or C terminus of said sequences or both contain substitutions and lack 1-10 amino acids at the N or C terminus and wherein said variants in the context of said protein retain the ability to bind receptor for which at least one of said  $\beta$ -subunits is a ligand.

6. (Amended) A pharmaceutical composition which comprises the protein of claim 1 in admixture with a [suitable pharmaceutical] pharmaceutically acceptable excipient.

Claim 12, line 1, delete "12" and insert -- 11 -- therefor.

Claim 13, line 1, delete "12" and insert -- 11 -- therefor.

Claim 14, line 1, delete "13" and insert -- 12 -- therefor.

Claim 15, line 3, delete "14" and insert -- 13 -- therefor.

Claim 16, line 3, delete "15" and insert -- 14 -- therefor.

✓  
Please add the following claims 17-27:

15. 17. The protein of claim 1 which is of the formula  $\beta^1$ -(linker<sup>1</sup>)<sub>m</sub>- $\alpha$ -(linker<sup>2</sup>)<sub>n</sub>- $\beta^2$  (1).

16. 18. The protein of claim 17 wherein said  $\beta$  and  $\alpha$  subunits are linked in head-to-tail configuration.

17. 19. The protein of claim 18 wherein one of m and n is 0 and the other is 1 and wherein the linker is CTP.

18. 20. The protein of claim 19 wherein m is 0, n is 1 and linker<sup>2</sup> is CTP.

19. 21. The protein of claim 20 which is CG $\beta$ - $\alpha$ -CTP-FSH $\beta$ .

20. 22. The protein of claim 1 which is of the formula  $\beta^1$ -(linker<sup>1</sup>)<sub>m</sub>- $\beta^2$ -(linker<sup>2</sup>)<sub>n</sub>- $\alpha$  (2).